

ChemComm

Accepted Manuscript



This article can be cited before page numbers have been issued, to do this please use: M. Greaney, V. Bhat, P. Duspara, S. Seo and N. S. B. Abu Bakar, *Chem. Commun.*, 2015, DOI: 10.1039/C4CC09987G.



This is an *Accepted Manuscript*, which has been through the Royal Society of Chemistry peer review process and has been accepted for publication.

Accepted Manuscripts are published online shortly after acceptance, before technical editing, formatting and proof reading. Using this free service, authors can make their results available to the community, in citable form, before we publish the edited article. We will replace this *Accepted Manuscript* with the edited and formatted *Advance Article* as soon as it is available.

You can find more information about *Accepted Manuscripts* in the [Information for Authors](#).

Please note that technical editing may introduce minor changes to the text and/or graphics, which may alter content. The journal's standard [Terms & Conditions](#) and the [Ethical guidelines](#) still apply. In no event shall the Royal Society of Chemistry be held responsible for any errors or omissions in this *Accepted Manuscript* or any consequences arising from the use of any information it contains.

Cite this: DOI: 10.1039/c0xx00000x

www.rsc.org/xxxxxx

COMMUNICATION

Visible light promoted thiol-ene reactions using titanium dioxide

Venugopal T. Bhat, Petar A. Duspara, Sangwon Seo, Nor Syazwani Binti Abu Bakar and Michael F. Greaney*

Received (in XXX, XXX) Xth XXXXXXXXX 20XX, Accepted Xth XXXXXXXXX 20XX

DOI: 10.1039/b000000x

The radical addition of thiols to alkenes is reported under photoredox conditions, using visible light and TiO₂ as a cheap and readily available photocatalyst.

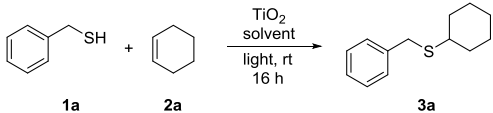
Recent developments in visible light photoredox catalysis (PRC) have substantially expanded the scope and application of organic photochemistry.¹ Using a domestic lightbulb as the light source, rather than specialist UV equipment, the process uses a photo-active catalyst to mediate electron transfer between reactants, resulting in novel transformations that frequently proceed under mild conditions. The vast majority of PRC in organic synthesis to date is based on two classes of catalyst; transition metal complexes, such as the archetypal Ru(bpy)₃Cl₂ and Ir(ppy)₃ systems, or organic dyes such as methylene blue or eosin-Y. The Noble metals of the first class are expensive and a limited resource, but are highly optimisable in terms of ligand-metal tuning. Organic dyes lack this particular quality, but may be tuned via conventional synthesis and are usually less expensive. We were interested in exploiting a third class of photoredox catalyst; semi-conducting metal oxides such as titania (TiO₂). With 4 million tons of titania being produced per annum, the material is cheap, readily accessible, and non-toxic (commonly ingested as food additive E 171), as well as offering the process benefits of heterogeneous catalysis such as simple removal by filtration.² The UV photoredox properties of TiO₂ have been extensively investigated and applied on an industrial scale to areas such as water treatment and photovoltaics. Applications to organic synthesis, however, remain underdeveloped, particularly in the area of visible light photochemistry.^{3,4} We were interested in developing a titania-mediated PRC reaction using visible light, and chose to investigate the thiol-ene reaction for this purpose.

The addition of thiols to alkenes is a radical reaction that is typically initiated with UV light.⁵ Although an old reaction,⁶ its reliability, substrate scope, and compatibility has led to renewed interest in the context of click chemistry, where the facile C-S bond formation can be used as a powerful ligation strategy in biological and materials science.⁷ Indeed, the largest field of application for thiol-ene chemistry is currently in polymer science, where thiols are cross-linked with alkenes under UV curing conditions, producing polymers with very uniform step-growth properties.⁸ Recent literature reports have established PRC for thiol-ene addition using both Ru(II) and eosin-Y

photocatalysis, providing an important frame of reference for our inchoate titania process.^{9,10,11}

We were aware at the outset that titania visible light PRC would likely require a nanoparticulate form of the catalyst. Unmodified titania absorbs very weakly in the visible region, but there is evidence that substrate binding to the surface can shift the absorption and enable electron transfer.^{4a,c,d,e} Rueping and co-workers, for example, have recently demonstrated visible light photoredox activity in dehydrogenative coupling reactions using commercially available titania P25 nanoparticles,^{4d,e} and we chose to begin our studies using this material (aeroxide® from Sigma-Aldrich). Using the addition of benzyl mercaptan (**1a**) to cyclohexene (**2a**) as a test reaction, we were pleased to observe very high conversions into adduct **3a** using one equivalent of TiO₂ in MeCN under irradiation with a domestic light bulb (Table 1). The reaction was also viable in DCM (entry 2), under neat conditions (entry 3), and using blue LEDs as an alternative light source (entry 4).

Table 1. Titania PRC of the thiol-ene reaction^a



entry	TiO ₂ (equiv.)	solvent	light source	yield ^b (%)
1	1.0	MeCN	20W light	98 (96 ^c)
2	1.0	DCM	20W light	96
3	1.0	-	20W light	97
4	1.0	MeCN	Blue LED	80
5 ^d	1.0	MeCN	no light	0
6 ^e	0	MeCN	20W light	20
7	0	MeCN	Blue LED	0
8	0.5	MeCN	20W light	88
9	0.1	MeCN	20W light	82
10 ^e	0.1	MeCN	20W light	92
11 ^f	1.0	MeCN	20W light	50
12 ^g	1.0	MeCN	20W light	84

^a Cyclohexene (1.0 mmol), benzyl mercaptan (4.0 mmol), titania (TiO₂ P25) and solvent (1.0 mL) at room temperature for 16 h; Light source placed 10–15 cm away from reaction vials. ^b Yields determined by ¹H NMR using nitromethane as the internal standard. ^c Isolated yields. ^d Foil-wrapped flask for 16 h. ^e Reaction for 40 h. ^f Using degassed MeCN under N₂ atmosphere. ^g Using 2.5 mmol of benzyl mercaptan.

Control experiments established that titania in the dark gave no reaction (entry 5), but there was a small background reaction with ambient light in the absence of titania (20% conversion). We were pleased to see that the titania loading could be reduced to catalytic quantities with little drop in overall yield (entries 8-10), and we established that oxygen was beneficial for the reaction. Use of an inert atmosphere and de-gassed solvent halved the efficiency of the process (entry 11). This was an expected requirement, as an oxidative quencher is needed to balance charge and prevent hole-electron recombination in the titania photocatalyst (*vide infra*).

With a titania PRC system in place, we moved on to investigate substrate scope, beginning with the alkene partner.

TABLE 2. Scope of alkene coupling partners^a

entry	alkene	product	yield ^b (%)
1	2b (allyl methyl ether)	3b	85 (9)
2	2c (allyl acetate)	3c	75 (61)
3	2d (3-methyl-2-propen-1-ol)	3d	78 (12)
4	2e (3-methyl-2-propen-1-ol)	3e	89 (68)
5	2f (allyl trimethylsilane)	3f	97 (31)
6	2g (acrylonitrile)	3g	43 (0)
7	2h (vinyl acetate)	3h	80 (0)
8	2i (styrene)	3i	98 (90)
9	2j (4-methylstyrene)	3j	98 (93)
10	2k (4-chlorostyrene)	3k	95 (30)

^a Alkene (1.0 mmol), benzyl mercaptan (4.0 mmol), titania (TiO₂ P25, 1.0 mmol) and MeCN (1.0 mL) at room temperature for 16 h; Light source placed 10-15 cm away from reaction vials. ^b Isolated yields; numbers in parentheses are the yield obtained under standard reaction conditions where titanium dioxide is absent.

We were pleased to find that a variety of primary and 1,1-disubstituted substrates underwent successful thiol-ene reaction with benzyl mercaptan (Table 2). The reaction was tolerant of a variety of functional groups (ester, alcohol, silane, nitrile, halogen) giving the thiol-ene adducts **3** in generally high yields. Substitution was tolerated at the 2- and 3-positions of the alkene (entries 3 and 4), but 1,2-disubstitution was restricted to cyclic alkenes. Vinyl acetate and acrylonitrile were both productive, interestingly undergoing no reaction at all in the absence of PRC (entries 6 and 7). More reactive alkenes such as styrenes gave very high yields of the thiol-ene products, but background reaction was high for **2i** and **2j**.

TABLE 3. Catalytic thiol-ene – thiol scope^{a,b}

entry	thiol	product	yield ^c (%)
1	1b (1-octanethiol)	3l	86 (0)
2	1c (6-mercaptohexan-1-ol)	3m	79 (8)
3	1d (4-mercapto-2-methylpentan-3-one)	3n	87 (30)
4	1e (benzenethiol)	3o	61 (14)
5	1f (cyclohexanethiol)	3p	30 (5)
6 ^d	1g (2-methyl-2-propanethiol)	-	0
7	1h (1,2-ethanedithiol)	3q	68 (0)
8	1h (1,2-ethanedithiol)	3r	73 (0)
9	1i (1,3-ethanedithiol)	3s	70 (0)
10 ^d	1h (1,2-ethanedithiol)	3t	57 (0)

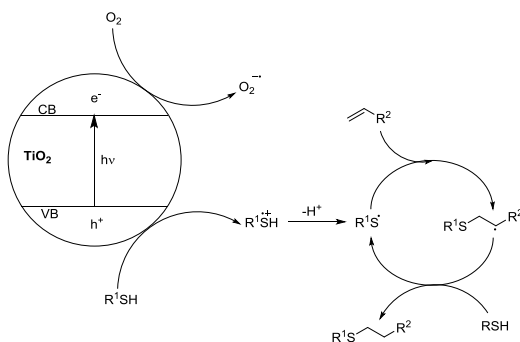
^a Thiol (4.0 mmol), vinyl acetate (1.0 mmol), titania (TiO₂ P25, 0.1 mmol) and MeCN (1.0 mL) at room temperature for 16 - 40 h; Light source placed 10-15 cm away from reaction vials. ^b Dithiol (1 mmol), styrene (4 mmol) and conditions as previous. ^c Isolated yields; numbers in parentheses are the yield obtained under standard reaction conditions where titanium dioxide is absent. ^d 1 equiv of titania used.

Thiol scope was initially trialled (Table 3) using catalytic amounts of titania (0.1 equiv) and vinyl acetate (**2h**), an alkene with zero background reaction for the addition of benzyl

mercaptan (Table 1, entry 7). We were pleased to find that the catalytic protocol was successful, with primary alkyl thiols (entries 1-3) and thiophenols (entry 4) undergoing thiol-ene reaction under the longer reaction time of 40 h. Increasing steric hindrance around the thiol was deleterious, with cyclohexyl thiol reacting in moderate yield (entry 5) and *tert*-butyl thiol undergoing no reaction, even with stoichiometric titania (entry 6). We could extend the reaction to a double thiol-ene reaction, with ethane and propane dithiol undergoing addition to styrenes in good yield (entries 7-9). *p*-Methoxystyrene proved less reactive, and required stoichiometric quantities of titania to yield **3t** in an acceptable timeframe.

A mechanism for the reaction is outlined in Scheme 1, based upon the photo-excitation of electrons to the valence band of the titania catalyst. The resultant holes are reductively quenched by the thiol, generating a thiyl radical cation, which can lose a proton to give a thiyl radical. Oxygen acts as a sacrificial electron acceptor, enhancing reaction efficiency by reducing hole-electron recombination in the titania. The thiyl radical can then initiate the thiol-ene cycle through addition to an alkene and generation of an alkyl radical, which propagates the reaction by abstracting a hydrogen atom from the thiol starting material.

Scheme 1. Mechanistic pathway for titania thiol-ene reaction



In conclusion, we have developed a titania photoredox system for the thiol-ene reaction that uses visible light. The photocatalyst is cheap, robust, readily available, and easily processed out of the reaction for re-use. Applications of this method to the broad remit of thiol-ene chemistry are ongoing in our laboratory.

We thank the University of Manchester and the EPSRC for funding.

Notes and references

^a School of Chemistry, University of Manchester, Oxford Rd, Manchester M13 9PL, UK; E-mail: michaelgreaney@manchester.ac.uk

† Electronic Supplementary Information (ESI) available: Synthesis and characterisation data for all new compounds. See DOI: 10.1039/b000000x/

¹ Recent reviews: a) J. M. R. Narayanan and C. R. J. Stephenson, *Chem. Soc. Rev.*, 2011, **40**, 102-113; b) C. K. Prier, D. A. Rankic and D. W. C. MacMillan, *Chem. Rev.*, 2013, **113**, 5322-5363; c) S. Fukuzumi and K. Ohkubo, *Org. Biomol. Chem.*, 2014, **12**, 6059-6071; d) D. P. Hari and B. König, *Chem. Commun.*, 2014, **50**, 6688-6699; e) M. N. Hopkinson, B. Sahoo, J. L. Li and F. Glorius, *Chem. Eur. J.*, 2014, **20**, 3874-3886; f) D. A. Nicewicz and T. M. Nguyen, *ACS Catal.*, 2014, **4**, 355-360.

- ² C. Chen, W. Ma and J. Zhao, *Chem. Soc. Rev.*, 2010, **39**, 4206-4219; b) P. Roy, S. Berger and P. Schmuki, *Angew. Chem. Int. Ed.*, 2011, **50**, 2904-2939; c) B. O'Regan and M. Grätzel, *Nature*, 1991, **353**, 737-740.
- ³ Selected examples of UV titania photoredox catalysis: a) C. M. Wang and T. E. Mallouk, *J. Am. Chem. Soc.*, 1990, **112**, 2016-2018; b) C. W. Lai and T. E. Mallouk, *Chem. Commun.*, 1993, 1359-1361; c) S. Higashida, A. Harada, R. Kawakatsu, N. Fujiwara and M. Matsumura, *Chem. Commun.*, 2006, 2804-2806; d) X. J. Lang, H. W. Ji, C. C. Chen, W. H. Ma and J. C. Zhao, *Angew. Chem. Int. Ed.*, 2011, **50**, 3934-3937; e) D. W. Manley, R. T. McBurney, P. Miller, R. F. Howe, S. Rhydderch and J. C. Walton, *J. Am. Chem. Soc.*, 2012, **134**, 13580-13583; f) D. W. Manley, R. T. McBurney, P. Miller and J. C. Walton, *J. Org. Chem.*, 2014, **79**, 1386-1398; g) D. W. Manley and J. C. Walton, *Org. Lett.*, 2014, **16**, 5394-5397.
- ⁴ With visible light: a) F. Parrino, A. Ramakrishnan and H. Kisch, *Angew. Chem. Int. Ed.*, 2008, **47**, 7107-7109; b) M. Cherevatskaya, M. Neumann, S. Fuldner, C. Harlander, S. Kummel, S. Dankesreiter, A. Pfützner, K. Zeitler and B. König, *Angew. Chem. Int. Ed.*, 2012, **51**, 4062-4066; c) X. Lang, W. Ma, Y. Zhao, C. Chen, H. Ji and J. Zhao, *Chem. Eur. J.*, 2012, **18**, 2624-2631; d) M. Rueping, J. Zoller, D. C. Fabry, K. Poschorny, R. M. Koenigs, T. E. Weirich and J. Mayer, *Chem. Eur. J.*, 2012, **18**, 3478-3481; e) C. Vila and M. Rueping, *Green Chem.*, 2013, **15**, 2056-2059; f) C. D. McTiernan, S. P. Pitre, H. Ismaili and J. C. Scaiano, *Adv. Synth. Catal.*, 2014, **356**, 2819-2824.
- ⁵ F. Dénès, M. Pichowicz, G. Povie, and P. Renaud, *Chem. Rev.*, 2014, **114**, 2587-2693.
- ⁶ T. Posner, *Ber. Dtsch. Chem. Ges.*, 1905, **38**, 646-657.
- ⁷ C. E. Hoyle, C. N. Bowman, *Angew. Chem., Int. Ed.*, 2010, **49**, 1540-1573.
- ⁸ a) A. B. Lowe, *Polymer Chem.*, 2014, **5**, 4820-4870; b) N. Brummelhuis, C. Diehl, and H. Schlaad, *Macromolecules*, 2008, **41**, 9946-9947; and references therein.
- ⁹ a) E. L. Tyson, M. S. Ament, and T. P. Yoon, *J. Org. Chem.*, 2013, **78**, 2046-2050; b) E. L. Tyson, Z. L. Niemeyer, and T. P. Yoon, *J. Org. Chem.*, 2014, **79**, 1427-1436.
- ¹⁰ M. H. Keylor, J. E. Park, C.-J. Wallentin, and C. R. J. Stephenson, *Tetrahedron*, 2014, **70**, 4264-4269.
- ¹¹ a) C. A. DeForest and K. S. Anseth, *Nat. Chem.*, 2011, **3**, 925-931; b) C. A. DeForest and K. S. Anseth, *Angew. Chem. Int. Ed.*, 2012, **51**, 1816-1819; c) H. Shih and C. C. Lin, *Macromol. Rapid Commun.*, 2013, **34**, 269-273; d) H. Shih, A. K. Fraser and C. C. Lin, *ACS Appl. Mater. Interfaces*, 2013, **5**, 1673-1680.